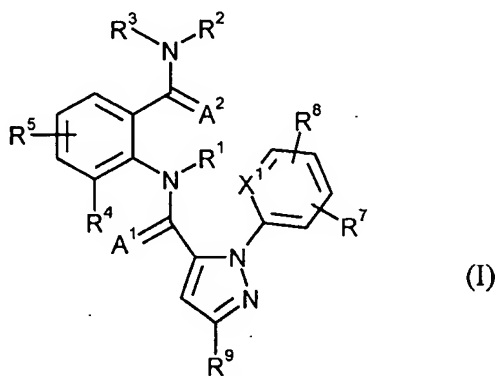


Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application.

1. (Currently amended) A composition comprising a synergistically effective active compound combination of at least one anthranilamide of the formula (I)



in which

A¹ and A² independently of one another represent oxygen or sulfur,

X¹ represents N or CR¹⁰,

R¹ represents hydrogen or represents C₁-C₆-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl or C₃-C₆-cycloalkyl, each of which is optionally mono- or polysubstituted, where the substituents independently of one another may be selected from the group consisting of R⁶, halogen, cyano, nitro, hydroxyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₂-C₄-alkoxycarbonyl, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₃-C₆-cycloalkylamino, (C₁-C₄-alkyl)-C₃-C₆-cycloalkylamino and R¹¹,

R² represents hydrogen, C₁-C₆-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl, C₃-C₆-cycloalkyl, C₁-C₄-alkoxy, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₃-C₆-cycloalkylamino, C₂-C₆-alkoxycarbonyl or C₂-C₆-alkylcarbonyl,

R³ represents hydrogen, R¹¹ or represents C₁-C₆-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl, C₃-C₆-cycloalkyl, each of which is optionally mono- or polysubstituted, where the substituents independently of one another may be selected from the group consisting of R⁶, halogen, cyano, nitro, hydroxyl, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₂-C₆-alkoxycarbonyl, C₂-C₆-alkylcarbonyl, C₃-C₆-trialkylsilyl, R¹¹, phenyl, phenoxy and a 5- or 6-membered heteroaromatic ring, where each phenyl, phenoxy and 5- or 6-membered heteroaromatic ring may optionally be substituted and where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R¹², or

R² and R³ may be attached to one another and form the ring M,

R⁴ represents hydrogen, C₁-C₆-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl, C₃-C₆-cycloalkyl, C₁-C₆-haloalkyl, C₂-C₆-haloalkenyl, C₂-C₆-haloalkynyl, C₃-C₆-halocycloalkyl, halogen, cyano, nitro, hydroxyl, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-haloalkylthio, C₁-C₄-haloalkylsulfinyl, C₁-C₄-haloalkylsulfonyl, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₃-C₆-cycloalkylamino, C₃-C₆-trialkylsilyl or represents phenyl, benzyl or phenoxy, each of which is optionally mono- or polysubstituted, where the substituents independently of one another may be

selected from the group consisting of C₁-C₄-alkyl, C₂-C₄-alkenyl, C₂-C₄-alkynyl, C₃-C₆-cycloalkyl, C₁-C₄-haloalkyl, C₂-C₄-haloalkenyl, C₂-C₄-haloalkynyl, C₃-C₆-halocycloalkyl, halogen, cyano, nitro, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₃-C₆-cycloalkylamino, C₃-C₆-(alkyl)cycloalkylamino, C₂-C₄-alkylcarbonyl, C₂-C₆-alkoxycarbonyl, C₂-C₆-alkylaminocarbonyl, C₃-C₈-dialkylaminocarbonyl and C₃-C₆-trialkylsilyl,

R⁵ and R⁸ in each case independently of one another represent hydrogen, halogen or represent in each case optionally substituted C₁-C₄-alkyl, C₁-C₄-haloalkyl, R¹², G, J, -OJ, -OG, -S(O)_p-J, -S(O)_p-G, -S(O)_p-phenyl, where the substituents independently of one another may be selected from one to three radicals W or from the group consisting of R¹², C₁-C₁₀-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl, C₁-C₄-alkoxy and C₁-C₄-alkylthio, where each substituent may be substituted by one or more substituents independently of one another selected from the group consisting of G, J, R⁶, halogen, cyano, nitro, amino, hydroxyl, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-haloalkylthio, C₁-C₄-haloalkylsulfinyl, C₁-C₄-haloalkylsulfonyl, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₃-C₆-trialkylsilyl, phenyl and phenoxy, where each phenyl or phenoxy ring may optionally be substituted and where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R¹²,

- G in each case independently of one another represents a 5- or 6-membered non-aromatic carbocyclic or heterocyclic ring which may optionally contain one or two ring members from the group consisting of C(=O), SO and S(=O)₂ and which may optionally be substituted by one to four substituents independently of one another selected from the group consisting of C₁-C₂-alkyl, halogen, cyano, nitro and C₁-C₂-alkoxy, or independently of one another represents C₂-C₆-alkenyl, C₂-C₆-alkynyl, C₃-C₇-cycloalkyl, (cyano)-C₃-C₇-cycloalkyl, (C₁-C₄-alkyl)-C₃-C₆-cycloalkyl, (C₃-C₆-cycloalkyl)-C₁-C₄-alkyl, where each cycloalkyl, (alkyl)cycloalkyl and (cycloalkyl)alkyl may optionally be substituted by one or more halogen atoms,
- J in each case independently of one another represents an optionally substituted 5- or 6-membered heteroaromatic ring, where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R¹²,
- R⁶ independently of one another represents -C(=E¹)R¹⁹, -LC(=E¹)R¹⁹, -C(=E¹)LR¹⁹, -LC(=E¹)LR¹⁹, -OP(=Q)(OR¹⁹)₂, -SO₂LR¹⁸ or -LSO₂LR¹⁹, where each E¹ independently of one another represents O, S, N-R¹⁵, N-OR¹⁵, N-N(R¹⁵)₂, N-S=O, N-CN or N-NO₂,
- R⁷ represents hydrogen, C₁-C₄-alkyl, C₁-C₄-haloalkyl, halogen, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-haloalkylthio, C₁-C₄-haloalkylsulfinyl, C₁-C₄-haloalkylsulfonyl,

- R⁹ represents C₁-C₄-haloalkyl, C₁-C₄-haloalkoxy, C₁-C₄-haloalkylsulfinyl or halogen,
- R¹⁰ represents hydrogen, C₁-C₄-alkyl, C₁-C₄-haloalkyl, halogen, cyano or C₁-C₄-haloalkoxy,
- R¹¹ in each case independently of one another represents in each case optionally mono- to trisubstituted C₁-C₆-alkylthio, C₁-C₆-alkylsulfenyl, C₁-C₆-haloalkylthio, C₁-C₆-haloalkylsulfenyl, phenylthio or phenylsulfenyl, where the substituents independently of one another may be selected from the list W, -S(O)_nN(R¹⁶)₂, -C(=O)R¹³, -L(C=O)R¹⁴, -S(C=O)LR¹⁴, -C(=O)LR¹³, -S(O)_nNR¹³C(=O)R¹³, -S(O)_nNR¹³C(=O)LR¹⁴ or -S(O)_nNR¹³S(O)₂LR¹⁴,
- L in each case independently of one another represents O, NR¹⁸ or S,
- R¹² in each case independently of one another represents -B(OR¹⁷)₂, amino, SH, thiocyanato, C₃-C₈-trialkylsilyloxy, C₁-C₄-alkyl disulfides, -SF₅, -C(=E¹)R¹⁹, -LC(=E¹)R¹⁹, -C(=E¹)LR¹⁹, -LC(=E¹)LR¹⁹, -OP(=Q)(OR¹⁹)₂, -SO₂LR¹⁹ or -LSO₂LR¹⁹,
- Q represents O or S,
- R¹³ in each case independently of one another represents hydrogen or represents in each case optionally mono- or polysubstituted C₁-C₆-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl or C₃-C₆-cycloalkyl, where the substituents independently of one another may be selected from the group consisting of R⁶, halogen, cyano, nitro, hydroxyl, C₁-C₄-alkoxy, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-

alkylamino, C₂-C₈-dialkylamino, C₃-C₆-cycloalkylamino or (C₁-C₄-alkyl)-C₃-C₆-cycloalkylamino,

R¹⁴ in each case independently of one another represents in each case optionally mono- or polysubstituted C₁-C₂₀-alkyl, C₂-C₂₀-alkenyl, C₂-C₂₀-alkynyl or C₃-C₆-cycloalkyl, where the substituents independently of one another may be selected from the group consisting of R⁶, halogen, cyano, nitro, hydroxyl, C₁-C₄-alkoxy, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₃-C₆-cycloalkylamino and (C₁-C₄-alkyl)-C₃-C₆-cycloalkylamino or represent optionally substituted phenyl, where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R¹²,

R¹⁵ in each case independently of one another represents hydrogen or represents in each case optionally mono- or polysubstituted C₁-C₆-haloalkyl or C₁-C₆-alkyl, where the substituents independently of one another may be selected from the group consisting of cyano, nitro, hydroxyl, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-haloalkylthio, C₁-C₄-haloalkylsulfinyl, C₁-C₄-haloalkylsulfonyl, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₂-C₆-alkoxycarbonyl, C₂-C₆-alkylcarbonyl, C₃-C₆-trialkylsilyl and optionally substituted phenyl, where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R¹², or N(R¹⁵)₂ represents a cycle which forms the ring M,

- R^{16} represents C_1 - C_{12} -alkyl or C_1 - C_{12} -haloalkyl, or $N(R^{16})_2$ represents a cycle which forms the ring M,
- R^{17} in each case independently of one another represents hydrogen or C_1 - C_4 -alkyl, or $B(OR^{17})_2$ represents a ring in which the two oxygen atoms are attached via a chain having two to three carbon atoms which are optionally substituted by one or two substituents independently of one another selected from the group consisting of methyl and C_2 - C_6 -alkoxycarbonyl,
- R^{18} in each case independently of one another represents hydrogen, C_1 - C_6 -alkyl or C_1 - C_6 -haloalkyl, or $N(R^{13})(R^{18})$ represents a cycle which forms the ring M,
- R^{19} in each case independently of one another represents hydrogen or represents in each case mono- or polysubstituted C_1 - C_6 -alkyl, where the substituents independently of one another may be selected from the group consisting of cyano, nitro, hydroxyl, C_1 - C_4 -alkoxy, C_1 - C_4 -haloalkoxy, C_1 - C_4 -alkylthio, C_1 - C_4 -alkylsulfinyl, C_1 - C_4 -alkylsulfonyl, C_1 - C_4 -haloalkylthio, C_1 - C_4 -haloalkylsulfinyl, C_1 - C_4 -haloalkylsulfonyl, C_1 - C_4 -alkylamino, C_2 - C_8 -dialkylamino, CO_2H , C_2 - C_6 -alkoxycarbonyl, C_2 - C_6 -alkylcarbonyl, C_3 - C_6 -trialkylsilyl and optionally substituted phenyl, where the substituents independently of one another may be selected from one to three radicals W, C_1 - C_6 -haloalkyl, C_3 - C_6 -cycloalkyl or phenyl or pyridyl, each of which is optionally mono- to trisubstituted by W,
- M in each case represents an optionally mono- to tetrasubstituted ring which, in addition to the nitrogen atom attached to the substituent pair R^{13} and R^{18} , $(R^{15})_2$

or (R¹⁶)₂, contains two to six carbon atoms and optionally additionally a further nitrogen, sulfur or oxygen atom, where the substituents independently of one another may be selected from the group consisting of C₁-C₂-alkyl, halogen, cyano, nitro and C₁-C₂-alkoxy,

- W in each case independently of one another represents C₁-C₄-alkyl, C₂-C₄-alkenyl, C₂-C₄-alkynyl, C₃-C₆-cycloalkyl, C₁-C₄-haloalkyl, C₂-C₄-haloalkenyl, C₂-C₄-haloalkynyl, C₃-C₆-halocycloalkyl, halogen, cyano, nitro, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-alkylamino, C₂-C₈-dialkylamino, C₃-C₆-cycloalkylamino, (C₁-C₄-alkyl)-C₃-C₆-cycloalkylamino, C₂-C₄-alkylcarbonyl, C₂-C₆-alkoxycarbonyl, CO₂H, C₂-C₆-alkylaminocarbonyl, C₃-C₈-dialkylaminocarbonyl or C₃-C₆-trialkylsilyl,
- n in each case independently of one another represents 0 or 1,
- p in each case independently of one another represents 0, 1 or 2,

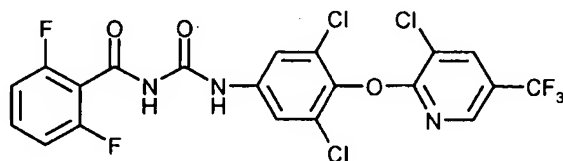
where, if (a) R⁵ represents hydrogen, C₁-C₆-alkyl, C₁-C₆-haloalkyl, C₂-C₆-haloalkenyl, C₂-C₆-haloalkynyl, C₁-C₄-haloalkoxy, C₁-C₄-haloalkylthio or halogen and (b) R⁸ represents hydrogen, C₁-C₆-alkyl, C₁-C₆-haloalkyl, C₂-C₆-haloalkenyl, C₂-C₆-haloalkynyl, C₁-C₄-haloalkoxy, C₁-C₄-haloalkylthio, halogen, C₂-C₄-alkylcarbonyl, C₂-C₆-alkoxycarbonyl, C₂-C₆-alkylaminocarbonyl or C₃-C₈ dialkylaminocarbonyl, (c) at least one substituent selected from the group consisting of R⁶, R¹¹ and R¹² if present and (d) if R¹² is not present, at least one of the radicals R⁶ and R¹¹ is different

from C₂-C₆-alkylcarbonyl, C₂-C₆-alkoxycarbonyl, C₂-C₆-alkylaminocarbonyl and C₃-C₈-dialkylaminocarbonyl, and
where the compound of the general formula (I) may also be an N-oxide or salt,

and at least one insecticidally active compound of group 2 below, selected from

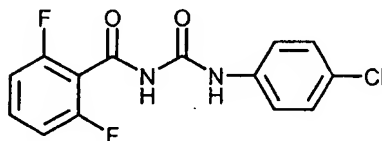
A) benzoylureas,

(2-1) chlorfluazuron



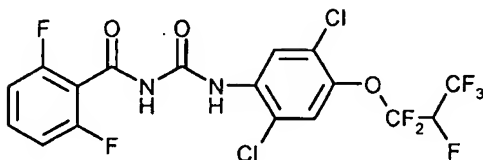
and/or

(2-2) diflubenzuron



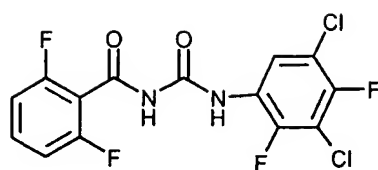
and/or

(2-3) lufenuron



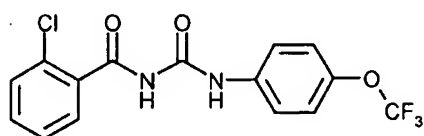
and/or

(2-4) teflubenzuron



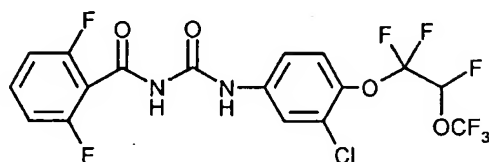
and/or

(2-5) triflumuron



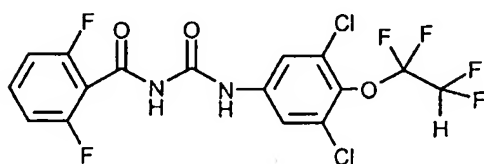
and/or

(2-6) novaluron



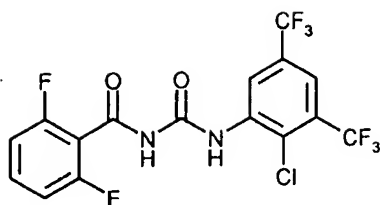
and/or

(2-7) hexaflumuron



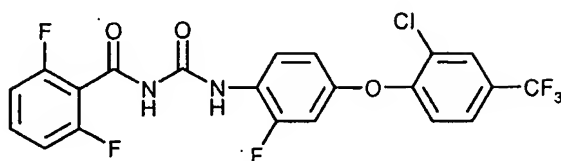
and/or

(2-8) bistrifluoron (DBI-3204)



and/or

(2-22) flufenoxuron



and/or

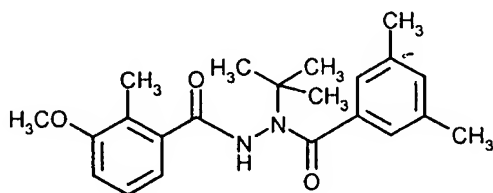
B) macrolides,

(2-9) emamectin

and/or

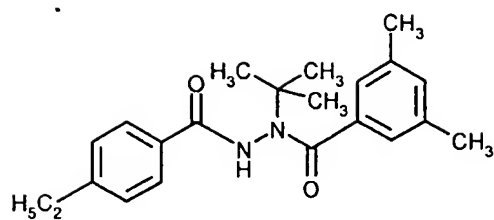
C) diacylhydrazines,

(2-10) methoxyfenozide



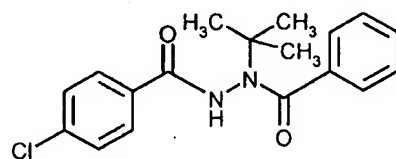
and/or

(2-11) tebufenozide



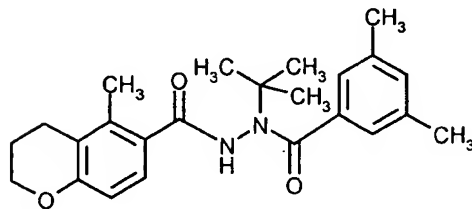
and/or

(2-12) halofenozide



and/or

(2-13) chromafenozide (ANS-118)



and/or

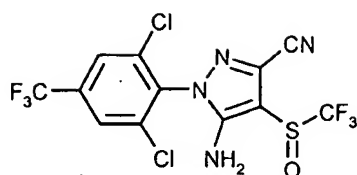
(2-14) *Trichogramma* spp.

and/or

(2-15) *Verticillium lecanii*

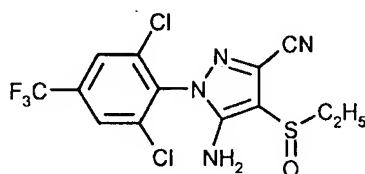
and/or

(2-16) fipronil



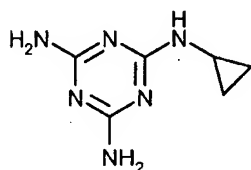
and/or

(2-17) ethiprole



and/or

(2-18) cyromazine

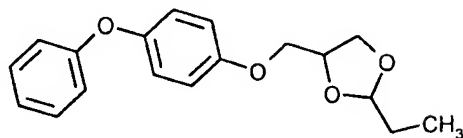


and/or

(2-19) azadirachtin

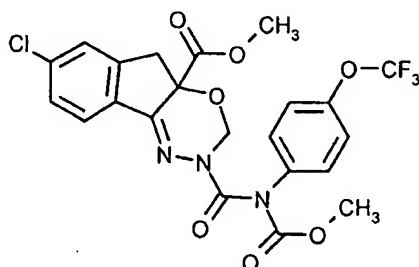
and/or

(2-20) diofenolan



and/or

(2-21) indoxacarb



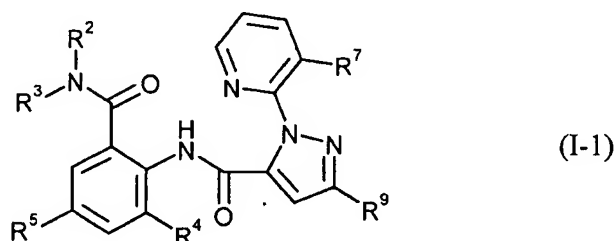
wherein the ratio of the at least one anthranilamide of formula (I) to the at least one active compound of group 2 is in a ratio of

<u>Formula (I) to chlorfluazuron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to diflubenzuron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to lufenuron</u>	<u>20 : 1 to 1 : 5</u>
<u>Formula (I) to teflubenzuron</u>	<u>20 : 1 to 1 : 5</u>
<u>Formula (I) to triflumuron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to novaluron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to hexaflumuron</u>	<u>20 : 1 to 1 : 5</u>
<u>Formula (I) to bistrifluoron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to flufenoxuron</u>	<u>50 : 1 to 1 : 5</u>
<u>Formula (I) to emamectin</u>	<u>50 : 1 to 1 : 5</u>
<u>Formula (I) to methoxyfenozide</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to tebufenozide</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to halofenozide</u>	<u>2 : 1 to 1 : 100</u>
<u>Formula (I) to chromafenozide</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to Trichogramma spp.</u>	<u>1000 g a.i./ha : 20 000 wasps/ha</u> <u>to</u> <u>10 g a.i./ha : 500 000 wasps/ha</u>
<u>Formula (I) to Verticillium lecanii</u>	<u>0.05 % a.i. : 0.05 % F^(*) to 0.001 %</u> <u>a.i. : 0.5 % F^(*)</u>
<u>Formula (I) to fipronil</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to ethiprole</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to cyromazine</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to azadirachtin</u>	<u>50 : 1 to 1 : 5</u>

<u>Formula (I) to diofenolan</u>	<u>100 : 1 to 1 : 2</u>
<u>Formula (I) to indoxacarb</u>	<u>50 : 1 to 1 : 5</u>

F(*) Formulation comprising 10^9 to 10^{10} spores/g.

2. (Original) The composition as claimed in claim 1 comprising at least one active compound from the group of the anthranilamides of the formula (I-1) in which



in which

- R^2 represents hydrogen or C_1 - C_6 -alkyl,
- R^3 represents C_1 - C_6 -alkyl which is optionally substituted by one R^6 ,
- R^4 represents C_1 - C_4 -alkyl, C_1 - C_2 -haloalkyl, C_1 - C_2 -haloalkoxy or halogen,
- R^5 represents hydrogen, C_1 - C_4 -alkyl, C_1 - C_2 -haloalkyl, C_1 - C_2 -haloalkoxy or halogen,
- R^6 represents $-C(=E^2)R^{19}$, $-LC(=E^2)R^{19}$, $-C(=E^2)LR^{19}$ or $-LC(=E^2)LR^{19}$, where each E^2 independently of one another represents O, S, $N-R^{15}$, $N-OR^{15}$, $N-N(R^{15})_2$, and each L independently of one another represents O or NR^{18} ,
- R^7 represents C_1 - C_4 -haloalkyl or halogen,
- R^9 represents C_1 - C_2 -haloalkyl, C_1 - C_2 -haloalkoxy, $S(O)_p$ - C_1 - C_2 -haloalkyl or halogen,

- R¹⁵ in each case independently of one another represents hydrogen or represents in each case optionally substituted C₁-C₆-haloalkyl or C₁-C₆-alkyl, where the substituents independently of one another may be selected from the group consisting of cyano, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-haloalkylthio, C₁-C₄-haloalkylsulfinyl and C₁-C₄-haloalkylsulfonyl,
- R¹⁸ in each case represents hydrogen or C₁-C₄-alkyl,
- R¹⁹ in each case independently of one another represents hydrogen or C₁-C₆-alkyl,
- p independently of one another represents 0, 1, 2.

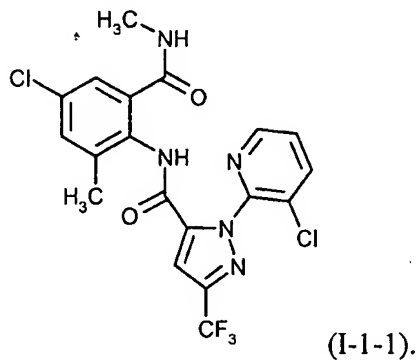
3. (Previously presented) The composition as claimed in claim 1 wherein the at least one active compound of group 2 is selected from the group consisting of
- (2-5) triflumuron
 - (2-22) flufenoxuron
 - (2-9) emamectin
 - (2-10) methoxyfenozide
 - (2-16) fipronil
 - (2-17) ethiprole and
 - (2-21) indoxacarb.
4. (Cancelled)

5. (Cancelled)
6. (Previously presented) A process for preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 1 is mixed with extenders and/or surfactants.
7. (Previously presented) A method for controlling animal pests, characterized in that synergistically effective mixtures as defined in claim 1 are allowed to act on animal pests and/or their habitat.
8. (Cancelled)
9. (Cancelled)
10. (Previously presented) A process for preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 2 is mixed with extenders and/or surfactants.
11. (Previously presented) A process for preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 3 is mixed with extenders and/or surfactants.

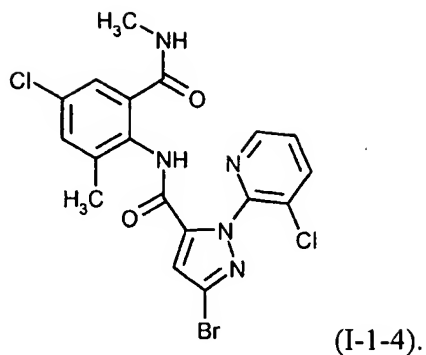
12. (Cancelled)
13. (Cancelled)
14. (Cancelled)
15. (Previously presented) A method for controlling animal pests, characterized in that synergistically effective mixtures as defined in claim 2 are allowed to act on animal pests and/or their habitat.
16. (Previously presented) A method for controlling animal pests, characterized in that synergistically effective mixtures as defined in claim 3 are allowed to act on animal pests and/or their habitat.
17. (Cancelled)
18. (Cancelled)
19. (Cancelled)
20. (Previously presented) The composition according to claim 1 wherein:
R² represents hydrogen or methyl,
R³ represents C₁-C₄-alkyl,

- R⁴ represents methyl, trifluoromethyl, trifluoromethoxy, fluorine, chlorine, bromine or iodine,
- R⁵ represents hydrogen, fluorine, chlorine, bromine, iodine, trifluoromethyl or trifluoromethoxy,
- R⁷ represents chlorine or bromine, and
- R⁹ represents trifluoromethyl, chlorine, bromine, difluoromethoxy or trifluoroethoxy.
21. (Previously presented) The composition according to claim 20 wherein the at least one active compound of group 2 is fipronil (2-16).
22. (Cancelled)
23. (Currently amended) The composition according to claim ~~[[22]]~~ 21 wherein the ratio of the at least one anthranilamide of formula (I) and fipronil (2-16) is from 5:1 to 1:5.
24. (Previously presented) The composition according to claim 23 wherein the ratio of the at least one anthranilamide of formula (I) and fipronil (2-16) is 1:5.
25. (Previously presented) The composition according to claim 20 wherein the at least one active compound of group 2 is ethiprole (2-17).
26. (Cancelled)

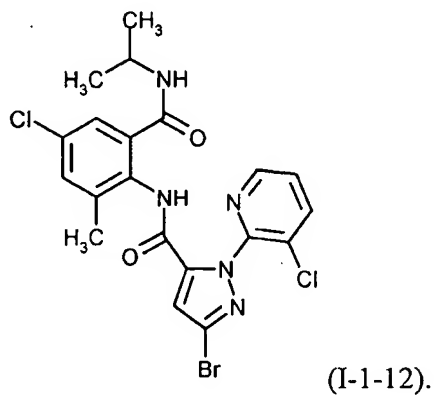
27. (Currently amended) The composition according to claim [[26]] 25 wherein the ratio of the at least one anthranilamide of formula (I) and ethiprole (2-17) is from 5:1 to 1:5.
28. (Previously presented) The composition according to claim 27 wherein the ratio of the at least one anthranilamide of formula (I) and ethiprole (2-17) is 1:5.
29. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



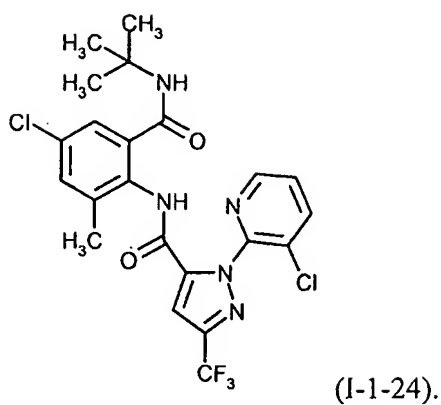
30. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



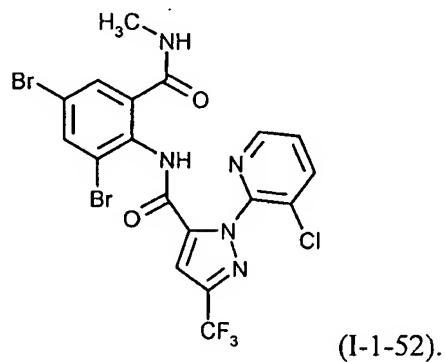
31. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



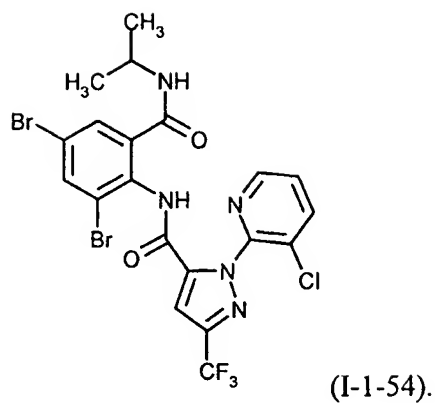
32. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



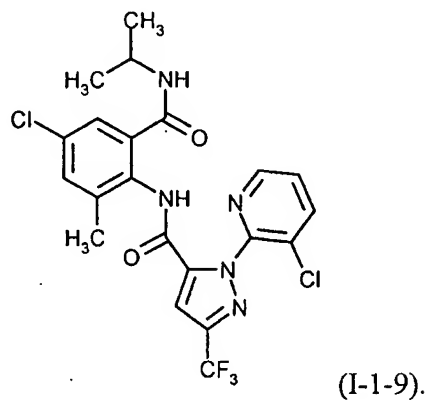
33. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



34. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



35. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



36. (Previously presented) The composition according to claim 35 wherein the at least one active compound of group 2 is fipronil (2-16).
37. (Previously presented) The composition according to claim 36 wherein the ratio of the compound (I-1-9) to fipronil (2-16) is 1:1.

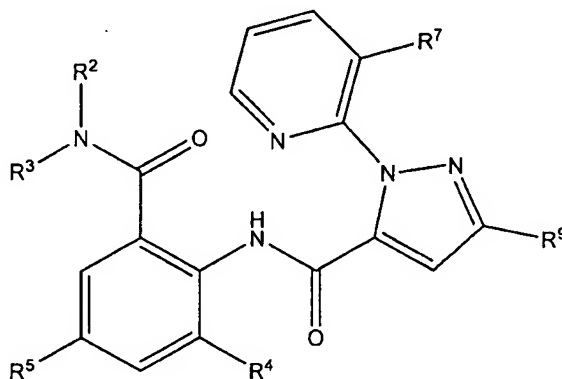
38. (Previously presented) The composition according to claim 35 wherein the at least one active compound of group 2 is ethiprole (2-17).
39. (Previously presented) The composition according to claim 38 wherein the ratio of the compound (I-1-9) to ethiprole (2-17) is 1:1.
40. (New) The composition according to claim 1, wherein the ratio of the at least one anthranilamide of formula (I) to the at least one active compound of group 2 is in a ratio of

Formula (I) to chlorfluazuron	5 : 1 to 1 : 5
Formula (I) to diflubenzuron	5 : 1 to 1 : 5
Formula (I) to lufenuron	10 : 1 to 1 : 2
Formula (I) to teflubenzuron	10 : 1 to 1 : 2
Formula (I) to triflumuron	5 : 1 to 1 : 5
Formula (I) to novaluron	5 : 1 to 1 : 5
Formula (I) to hexaflumuron	5 : 1 to 1 : 2
Formula (I) to bistrifluoron	5 : 1 to 1 : 5
Formula (I) to flufenoxuron	10 : 1 to 1 : 1
Formula (I) to emamectin	10 : 1 to 1 : 1
Formula (I) to methoxyfenozide	5 : 1 to 1 : 5
Formula (I) to tebufenozide	5 : 1 to 1 : 5
Formula (I) to halofenozide	1 : 1 to 1 : 30
Formula (I) to chromafenozide	5 : 1 to 1 : 5
Formula (I) to Trichogramma spp.	300 g a.i./ha : 50 000 wasps/ha to 50 g a.i./ha : 300 000 wasps/ha
Formula (I) to Verticillium lecanii	0.03 % a.i. : 0.1 % F ^(*) to 0.005 % a.i. : 0.2 % F ^(*)
Formula (I) to fipronil	5 : 1 to 1 : 5

Formula (I) to ethiprole	5 : 1 to 1 : 5
Formula (I) to cyromazine	5 : 1 to 1 : 5
Formula (I) to azadirachtin	10 : 1 to 1 : 1
Formula (I) to diofenolan	20 : 1 to 1 : 1
Formula (I) to indoxacarb	20 : 1 to 1 : 2

F(*) Formulation comprising 10^9 to 10^{10} spores/g.

41. (New) A process of preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 40 is mixed with extenders and/or surfactants.
42. (New) A method for controlling animal pests, characterized in that a synergistically effective mixture as defined in claim 40 is allowed to act on animal pests and/or their habitat.
43. (New) The composition according to claim 1, wherein said compound of formula (I) is one of:



Compound No.	R ²	R ³	R ⁴	R ⁵	R ⁷	R ⁹
I-1-1	H	Me	Me	Cl	Cl	CF ₃
I-1-4	H	Me	Me	Cl	Cl	Br
I-1-12	H	i-Pr	Me	Cl	Cl	Br
I-1-24	H	t-Bu	Me	Cl	Cl	CF ₃
I-1-52	H	Me	Br	Br	Cl	CF ₃
I-1-54	H	i-Pr	Br	Br	Cl	CF ₃

44. (New) The composition according to claim 43, wherein said at least one compound of group 2 is ethiprole or fipronil.
45. (New) The composition according to claim 44, wherein the ratio of the compound of formula (1) to ethiprole or fipronil is from 5:1 to 1:5.
46. (New) A process of preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 45 is mixed with extenders and/or surfactants.
47. (New) A method for controlling animal pests, characterized in that a synergistically effective mixture as defined in claim 45 is allowed to act on animal pests and/or their habitat.